

PATENT
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**SCREENING ASSAY FOR THE IDENTIFICATION OF
AGENTS WHICH ALTER EXPRESSION OF PTH-rP**

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BACKGROUND OF THE INVENTION

A1 A This application claims priority on provisional application U.S. Serial Number 60/025,215, filed August 23, 1996.

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Field of the Invention

The present invention relates generally to methods for screening compounds for their ability to affect the production of certain peptides which are destructive to those cells and to organisms, in some situations, but may be beneficial to those cells and organisms in other situations. More specifically, the invention relates to a method of identifying compounds from sources such as small molecule chemical libraries, peptide libraries and from natural product collections which inhibit the production of peptides related to bone destruction in patients with cancer, especially metastatic breast cancer. Furthermore, the invention as described specifically above may be used to identify compounds from said sources that stimulate the production of peptides related to bone formation and repair.

Related Art

Solid tumors such as breast cancer and lung cancer cause bone destruction by spreading to the bone marrow cavity, where tumor cells accumulate, grow and cause progressive bone destruction. These destructive bone lesions are known as osteolytic metastases and are responsible for complications in patients with cancer such as bone pain, pathologic fractures, hypercalcemia and nerve compression syndromes (Mundy, 1990).

One of the mechanisms by which breast cancer cells cause bone destruction is by producing the tumor peptide parathyroid hormone-related protein (PTH-rP). This protein was purified in 1987 simultaneously by several independent groups from human lung cancer (Moseley *et al.*, 1987), breast cancer (Burtis *et al.*, 1987) and renal cell carcinoma (Strewler *et al.*, 1987). Apparently, PTH-rP production in tumor cells is induced by high levels of TGF- β found in bone tissue.

Recent evidence supports the hypothesis that PTH-rP is important in the bone destruction mediated by breast cancer and other tumors which metastasize to bone. In breast cancer, PTH-rP was detected by immunohistochemical staining in 60% of 102 invasive breast cancers removed 5 from normocalcemic women, but not in normal breast tissue (Southby *et al.*, 1990). At least two other studies have confirmed these findings (Liapis *et al.*, 1993; Bundred *et al.*, 1992). By immunohistochemistry (Vargas *et al.*, 1992) and *in situ* hybridization (Powell *et al.*, 1991), PTH-rP was detected in 12 out of 13 breast cancer metastases in bone leading to the conclusion that 10 production of PTH-rP locally in bone may be responsible for the bone destruction associated with metastatic breast cancer.

Bundred and colleagues found positive immunohistochemical staining for PTH-rP in 56% of 155 primary breast tumors from normocalcemic women and PTH-rP correlated with development of bone metastases (Bundred *et al.*, 1992). PTH-rP expression was detected by 15 reverse transcriptase-PCR in 37 out of 38 primary breast cancers, and subsequent development of bone metastases was associated with greater PTH-rP expression (Bouizar *et al.*, 1993). Thus, PTH-rP expression appears to be common in breast cancers once they have metastasized to bone.

More recently, Yin *et al.* (1995) have studied the capacity of breast cancer cells to 20 produce PTH-rP in the bone microenvironment using established cultures of human breast cancer cell lines. These workers found that 4 of 8 established human breast cancer cell lines expressed detectable PTH-rP, and one of these PTH-rP secreting cell lines, MDA-MB-231, was studied in detail using an *in vivo* model of osteolytic metastases. In this model, cancer cells are inoculated 25 into the left ventricle of the heart of nude mice and the development of osteolytic metastases was followed by radiology and histology 3-6 weeks later. These workers found that there was increased PTH-rP concentrations in bone marrow plasma taken from the femurs of mice with osteolytic lesions, a 2.5-fold increase over corresponding plasma PTH-rP concentrations. This indicates local production of PTH-rP in the bone micro-environment. Moreover, when they 30 investigated the functional role of PTH-rP by inoculating the mice with neutralizing antibodies to PTH-rP, and compared them with mice receiving control IgG or no treatment, they found not

only decreased osteoclast number/mm² of tumor-bone interface and increased bone area, but also decreased tumor area in tumor bearing animals treated with PTH-rP antibodies compared with corresponding controls. These results show that tumor-produced PTH-rP may cause local bone destruction in human breast cancers which metastasize to bone. Moreover, they show that 5 neutralizing antibodies to PTH-rP reduce tumor burden in bone.

Clearly, PTH-rP production by tumor cells contributes in the pathogenic process of bone metastasis. It would, therefore, be highly desirable to identify compounds which inhibit PTH-rP production, thus halt progressive bone destruction and improve the quality of life of patients with 10 cancer. Thus, there is a need for techniques for detecting chemical compounds that decrease the production of PTH-rP by cancer cells.

PTH-rP influences a number of biological processes. Perhaps the most important is 15 endochondral bone formation. PTH-rP is known to be essential for normal cartilage differentiation and endochondral bone formation. Deletion of the PTH-rP gene by introducing the null mutant into the germ line results in mice with impaired skeletal development and enhanced endochondral ossification (Karaplis et al, 1994). PTH-rP is a negative regulator of 20 endochondral bone formation and terminal cartilage cell differentiation. This effect is possibly mediated by Indian Hedgehog protein which is made by pre- and early hypertrophic cells and stimulates production of PTH-rP in the perichondrium (Lanske et al, 1996; Vortkamp et al, 1996). Thus PTH-rP likely has an important role in normal endochondral bone formation and possibly also in fracture repair and therefore altering rates of PTH-rP expression may influence not just bone formation but also fracture repair, and such compounds could be administered to enhance the fracture repair process

25 PTH-rP also is produced by osteoblasts in the bone microenvironment. This has been demonstrated in osteoblast cultures as well as in organ cultures of rat long bones (Suda et al, 1996). PTH (and PTH-rP) have powerful anabolic effects on bone (Stewart 1996; Vickery et al, 1997), and therefore enhancing local production of PTH-rP in the bone microenvironment by 30 compounds which increase its transcription cause anabolic effects in this microenvironment.

SUMMARY OF THE INVENTION

The present invention addresses the foregoing need by providing a cell-based assay for identifying compounds which affect PTH-rP production. Specifically, the invention comprises 5 culturing a cell containing an expression vector comprising a DNA sequence encoding a PTH-rP promoter operatively linked to a reporter gene, namely firefly luciferase, under conditions which permit expression and quantitative assay of the reporter gene. The cultured cell is incubated with compounds suspected of possessing regulatory activity for production of PTH-rP. These inhibitory compounds are identified by their ability to modulate the expression of the reporter 10 gene and thereby affect the production of the assayable product of the reporter gene.

In a general embodiment, the present invention provides a method for screening a compound for its ability to affect PTH-rP production in mammalian cells. The method comprises the following steps:

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- (a) providing an expression construct comprising a PTH-rP promoter and a reporter gene, wherein the reporter gene is under transcriptional control of the promoter;
- (b) transfecting the mammalian cells with the expression construct
- (c) contacting the transfected cell with the compound; and
- (d) identifying a compound that regulates expression of the reporter gene from the 20 promoter.

In a preferred embodiment, the reporter gene is selected from the group consisting of firefly luciferase, chloramphenicol acetyl transferase, β -galactosidase, green fluorescent protein, 25 human growth hormone, alkaline phosphatase and β -glucuronidase. In a most preferred embodiment, the reporter gene is firefly luciferase.

In a further preferred embodiment, the promoter for PTH-rP is derived from the native PTH-rP promoter. Promoter sequences (P1, P2 and P3) are shown SEQ ID NO:1. A particularly

effective expression construct is the plasmid pGL3B-PTH-rP 1.1, containing the P2 and P3 promoters.

It is preferred that the present invention be used to screen a compound for its ability to 5 regulate PTH-rP production in human cells. A particularly useful cell population to use in screening for PTH-rP inhibition is human tumor cells. Most notably, the present invention is useful in screening compounds which affect PTH-rP production in breast cancer cells. A particularly useful breast cancer cell population in which to perform screening are MDA-MB-10 231 cells. Another especially preferred cell population for screening compounds which affect PTH-rP production are lung cancer cells. A particularly useful cell population in which to perform screening are RWGT2 cells. Another especially preferred cell population for screening compounds which affect production of PTH-rP are bone cells. A particularly useful cell population in which to perform screening are bone cells selected from the group consisting of 15 MC3T3-E1, MG-63, U2OS, UMR-106, ROS17/2.8 and SAOS-2.

In another general embodiment, the present invention provides compounds that affect 20 PTH-rP production in mammalian cells. This compound is identified by the method comprising the steps of:

25 (a) providing an expression construct comprising a PTH-rP promoter and a reporter gene, wherein the reporter gene is under transcriptional control of the promoter;

(b) transfecting the mammalian cells with the expression construct;

(c) contacting the transfected cell with the compound; and

(d) identifying a compound that regulates expression of the reporter gene from the promoter.

30 Preferably, the compound is identified from a small molecule chemical library, a peptide library, or from a collection of natural products. In a preferred embodiment, the compound is OSW_s1, OSW₃, OSW₆ and analogs thereof including but not limited to:

6-chloro-9-(tetrahydro-2-pyranyl) purine
6-methyl purine riboside
6-(β -hydroxyethylamino) purine
6-methylmercaptopurine
5 6-(2-tetrahydropyranyl) adenine
8-aminoguanosine
6-(α , β -dimethylallylamino) purine
6-(α , β -dimethylallylamino) purine riboside
cis-zeatin
10 6-(1-hydroxyguanidino) purine riboside
purine
purine riboside
aristeromycin
zeatin
15 n-benzyl-9-(2-tetrahydropyranyl) adenine
8-azaxanthine
8-aza-2,6 diamino purine
8-azaadenine
8-azaguanine
20 2-amino-6-(alpha-carboxyethyl)-mercaptopurine
2-amino-6-(1'-methyl-4'-nitro-5'-imidazoyl) mercaptopurine
2-amino-6-mercaptopurine riboside (6-mercaptopurine)
2-mercaptopurine (2-thiopurine)
2-thioxanthine (6-hydroxy-2-mercaptopurine)
25 6-mercaptopurine
6-mercaptopurine-2'-deoxyriboside
6-mercaptopurine riboside
6-mercaptopurine riboside 5'-phosphate (thioinosinic acid)
6-mercaptopurine arabinoside
6-thioxanthine (2-hydroxy-6-mercaptopurine)
30 2',3'-O-isopropylidene-6-mercaptopurine riboside

Finally, yet a third embodiment of the present invention provides a method of regulating
35 PTH-rP production in mammalian cells. This method comprises the step of contacting a cell
with a compound that affects PTH-rP production in the cell. Preferred compounds are OSWs1,
OSW2, OSW3, OSW4, and OSW6 and analogs thereof and dexamethasone.

Other objects, features and advantages of the present invention will become apparent
40 from the following detailed description. It should be understood, however, that the detailed

description and the specific examples, while indicating preferred embodiments of the invention, are given by way of illustration only, since various changes and modifications within the spirit and scope of the invention will become apparent to those skilled in the art from this detailed description.

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BRIEF DESCRIPTION OF THE DRAWINGS

The following drawings form part of the present specification and are included to further demonstrate certain aspects of the present invention. The invention may be better understood by 10 reference to one or more of these drawings in combination with the detailed description of specific embodiments presented herein:

15 FIG. 1 shows the compound 5-benzylidene hydrazino-1,2,34-tetrazole, known as OSW3.

20 FIG. 2 shows the compound 2-(2-carboxypyrid-2-yl)-4-methyl-4-propylimidazolidin-5-one(HCl), known as OSW2.

25 FIG. 3 shows the compound glaucarubolone, known as OSW4.

30 FIG. 4 shows the compound dexamethasone.

FIG. 5 shows the compound 6-thioguanine, known as OSW6.

FIG. 6 shows the compound acriflavinium hydrochloride, known as OSWs1.

35 FIG. 7 shows the effect of OSW3 in lowering whole blood calcium levels after evidence of hypercalcemia in nude mice.

40 FIG. 8A and FIG. 8B show the effect of OSW3 in reducing whole blood calcium levels prior to onset of hypercalcemia in nude mice.

FIG. 9 shows the effect of OSW3 in reducing serum PTH-rP levels prior to onset of hypercalcemia in nude mice.

5 FIG. 10A and FIG. 10B show the effect of OSW6 in lowering whole blood calcium levels after evidence of hypercalcemia in nude mice.

10 FIG. 11 shows the effect of OSW6 in reducing tumor volume after evidence of hypercalcemia in nude mice.

15 FIG. 12 shows the effect of OSW6 in lowering whole blood calcium levels prior to onset of hypercalcemia in nude mice.

20 FIG. 13A and FIG. 13B show the effect of OSW3 in reducing lesion number and lesion volume prior to onset of bone metastasis in nude mice.

FIG. 14A and FIG. 14B show the effect of dexamethasone in reducing lesion number and lesion volume prior to onset of bone metastasis in nude mice.

FIG. 15A and FIG. 15B shows the effect of OSWs1 in stimulating the PTH-rP promoter *in vitro*.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

As stated above, evidence suggests a role for PTH-rP in certain cancers along with bone formation and repair. The present invention involves a cell-based assay technique for identifying 5 and evaluating chemical compounds and agents which affect the production of PTH-rP, thereby identifying chemotherapeutic compounds for use in the treatment of cancer. This cell-based assay also is believed to work equally well in assessing compounds for their inhibition of PTH-rP production in a variety of cancers including lung, bone and breast cancers. Furthermore, this 10 cell-based assay is believed to work equally well in identifying compounds for their stimulation of PTH-rP production, thereby identifying therapeutic compounds for the use in bone repair (e.g., after bone fracture).

Specifically, cells are transfected with an expression vector comprising a DNA sequence 15 encoding a promoter region of PTH-rP operatively linked to a reporter gene encoding an assayable product. The cells are then cultured under conditions which permit expression of the assayable product. The PTH-rP promoter region is preferably cloned from genomic DNA but may be synthesized *de novo*. A preferred expression vector construct is the plasmid pGL3B-PTH-rP1.1 containing a PTH-rP promoter and the gene for firefly luciferase as the reporter gene.

After transfection with the expression vector, the cells are incubated with at least one 20 compound suspected of possessing regulatory activity for PTH-rP expression. Chemical agents and factors can be identified by their ability to modulate the expression of the reporter gene and thereby increase or decrease the production of the assayable product. Such chemical compounds are selected from small chemical libraries, peptide libraries, and/or collections of natural 25 products.

The present invention is distinguished from other techniques for identifying chemical 30 compounds, as it specifically identifies chemical compounds, agents, factors and other substances which affect PTH-rP production by cells. These agents are identified by their capacity to affect the activity of PTH-rP promoters. Decrease in activity of the promoters is

measured by a correspondent decrease in production of the reporter gene's product. Increase in activity of the promoters is measured by a correspondent increase in production of the reporter gene's product. Thus, decrease in the production of, for example, firefly luciferase, indicates that PTH-rP promoter activity is being suppressed by the compound being tested; an increase in the 5 production of firefly luciferase is indicative of stimulation of the PTH-rP promoter. The affect in production of the assaying product reflects the affect in PTH-rP that would occur in a cell treated with the compound.

Ultimately, when cancer patients are treated with chemical compounds shown to decrease 10 PTH-rP promoter activity, PTH-rP production by tumor cells will be inhibited, resulting in decreased bone destruction and impaired growth of the tumor in bone tissue. Therefore, compounds identified by this assay technique that decrease PTH-rP promoter activity can be used in the treatment of cancers which metastasize to bone and cause hypercalcemia, and other conditions where PTH-rP production is excessive and harmful.

15 When patients requiring bone formation or repair are treated with chemical compounds shown to increase PTH-rP promoter activity, PTH-rP production by osteoblasts will be stimulated, resulting in bone formation and repair. Therefore, compounds identified by this assay technique that increase PTH-rP promoter activity can be used in the treatment of bone 20 trauma, underdevelopment of the bone, and other conditions where PTH-rP production is desired.

A. PTH-rP PROMOTER

A technique often employed by those skilled in the art of protein production today is to 25 obtain a "recombinant" version of the protein, to express it in a recombinant cell and to obtain the protein from such cells. These techniques are based upon the "cloning" of a DNA molecule encoding the protein from a DNA library, *i.e.*, on obtaining a specific DNA molecule distinct from other portions of DNA. This can be achieved, for purposes of the present invention, by cloning of a genomic DNA molecule containing a PTH-rP promoter. Alternatively, having

knowledge of the PTH-rP promoter sequence, the promoter may be synthesized according to standard techniques.

5 The first step in a cloning procedure is the screening of an appropriate DNA library, such as, in the present case, a tumor-derived library. The screening procedure may be an expression screening protocol employing antibodies directed against the protein, or activity assays. Alternatively, screening may be based on the hybridization of oligonucleotide probes, designed from a consideration of portions of the amino acid sequence of the protein, or from the DNA sequences of genes encoding related proteins. The operation of such screening protocols are well 10 known to those of skill in the art and are described in detail in the scientific literature.

15 Three PTH-rP promoters are contained within a 4.3 kB genomic DNA upstream from the start site of the PTH-rP gene, located on chromosome 12. The three promoters are as follows: P1 is 5' of exon 1A (the upstream TATA element); P2 is 5' of exon 1C (the midregion GC rich element); P3 is 5' of exon 2 (downstream TATA element). Three isoforms of PTH-rP have been identified, which are 139, 141 and 173 amino acids in length. All three isoforms have the same 20 amino acid sequence through residue 139. The isoforms arise from alternative 3' splicing. Their relative secretory rates and their relative importance to normal and pathophysiology have yet to be fully elucidated. However, most, and possibly all tumors expressing PTH-rP express PTH-rP 1-141 driven by P3.

25 Constructs employed are pGL3B-PTHrP 1.1 and pGL4B-PTHrP 4.0, the former incorporating promoters P2 and P3 and the latter incorporating all three promoters. A 4345 bp fragment of the PTH-rP promoter region is set forth in SEQ ID NO:1.

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B. REPORTER GENES

30 A reporter gene is a gene which produces a product having a readily identifiable and assayable phenotype. The gene encoding firefly luciferase (Promega, Madison, WI) is particularly useful in the present invention because the gene produces an easily quantifiable, visually detectable enzyme. One skilled in the art will however recognize other useful reporter

genes which will work equally well in the present invention. Examples of such reporter genes include, but are not limited to, chloramphenicol acetyl transferase (Promega), β -galactosidase (Promega), green fluorescent protein (Clontech, Palo Alto, CA), human growth hormone (Amersham Life Science, Arlington Heights, IL), alkaline phosphatase (Clontech) and β -glucuronidase (Clontech).

C. EXPRESSION CONSTRUCTS

The expression constructs, commonly referred to as vectors, that can be utilized in the disclosed cell-based assay of the instant invention may vary considerably.

10 The vectors may be "standard" expression vectors, *i.e.*, plasmids that contain one or more effector genes and regulatory elements required for expression of the effector gene in cells. Plasmid expression vectors include any plasmid, cosmid or phage construct that is capable of supporting expression of encoded genes in mammalian cells, such as pUC or BluescriptTM plasmid series. Alternatively, these vectors may be more complex, such as the viral vectors discussed below.

15 The regulatory elements of an expression vector will comprise at least a promoter, in this case the PTH-rP promoter and a reporter gene (as discussed above), and also may include structures that assist in replication, such as origins of replication. In addition, almost all expression vectors contain multipurpose cloning regions that have numerous restriction enzyme sites. One also typically will include a polyadenylation signal to effect proper polyadenylation of the transcript. The nature of the polyadenylation signal is not believed to be crucial to the successful practice of the invention, and any such sequence may be employed. Examples include 20 SV40 and bovine growth hormone poly-A sites. Also contemplated as an element of the expression construct is a terminator. These elements can serve to enhance message levels and to minimize read through from the construct into other sequences. Finally, expression vectors typically have selectable markers, often in the form of antibiotic resistance genes, that permit 25 selection of cells that carry these vectors.

As stated above, in certain embodiments of the present invention, the expression construct comprises a virus or engineered construct derived from a viral genome. The ability of certain viruses to enter cells via receptor-mediated endocytosis and, in some cases, integrate into the host cell chromosomes, have made them attractive candidates for gene transfer in to 5 mammalian cells.

i. **Retroviruses**

The retroviruses are a group of single-stranded RNA viruses characterized by an ability to convert their RNA to double-stranded DNA in infected cells by a process of reverse-transcription 10 (Coffin, 1990). The resulting DNA then stably integrates into cellular chromosomes as a provirus and directs synthesis of viral proteins. The integration results in the retention of the viral gene sequences in the recipient cell and its descendants. The retroviral genome contains three genes - *gag*, *pol*, and *env* - that code for capsid proteins, polymerase enzyme, and envelope 15 components, respectively. A sequence found upstream from the *gag* gene, termed Ψ , functions as a signal for packaging of the genome into virions. Two long terminal repeat (LTR) sequences are present at the 5' and 3' ends of the viral genome. These contain strong promoter and enhancer sequences and are also required for integration in the host cell genome (Coffin, 1990).

In order to construct a retroviral vector, a nucleic acid encoding a PTH-rP promoter is 20 inserted into the viral genome in the place of certain viral sequences to produce a virus that is replication-defective. In order to produce virions, a packaging cell line containing the *gag*, *pol* and *env* genes but without the LTR and Ψ components is constructed (Mann *et al.*, 1983). When a recombinant plasmid containing a human cDNA, together with the retroviral LTR and Ψ 25 sequences is introduced into this cell line (by calcium phosphate precipitation for example), the Ψ sequence allows the RNA transcript of the recombinant plasmid to be packaged into viral particles, which are then secreted into the culture media (Nicolas and Rubenstein, 1988; Temin, 1986; Mann *et al.*, 1983). The media containing the recombinant retroviruses is then collected, optionally concentrated, and used for gene transfer. Retroviral vectors are able to infect a broad

variety of cell types. However, integration and stable expression require the division of host cells (Paskind *et al.*, 1975).

5 A novel approach designed to allow specific targeting of retrovirus vectors was recently developed based on the chemical modification of a retrovirus by the chemical addition of galactose residues to the viral envelope. This modification could permit the specific infection of cells such as hepatocytes via asialoglycoprotein receptors, should this be desired.

10 A different approach to targeting of recombinant retroviruses was designed in which biotinylated antibodies against a retroviral envelope protein and against a specific cell receptor were used. The antibodies were coupled via the biotin components by using streptavidin (Roux *et al.*, 1989). Using antibodies against major histocompatibility complex class I and class II antigens, the infection of a variety of human cells that bore those surface antigens was demonstrated with an ecotropic virus *in vitro* (Roux *et al.*, 1989).

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ii. Adenoviruses

20 Human adenoviruses are double-stranded DNA tumor viruses with genome sizes of approximate 36 kB (Tooze, 1981). As a model system for eukaryotic gene expression, adenoviruses have been widely studied and well characterized, which makes them an attractive system for development of adenovirus as a gene transfer system. This group of viruses is easy to grow and manipulate, and exhibit a broad host range *in vitro* and *in vivo*. In lytically infected cells, adenoviruses are capable of shutting off host protein synthesis, directing cellular machinery to synthesize large quantities of viral proteins, and producing copious amounts of virus.

25 The E1 region of the genome includes E1A and E1B which encode proteins responsible for transcription regulation of the viral genome, as well as a few cellular genes. E2 expression, including E2A and E2B, allows synthesis of viral replicative functions, *e.g.* DNA-binding protein, DNA polymerase, and a terminal protein that primes replication. E3 gene products prevent cytolysis by cytotoxic T cells and tumor necrosis factor and appear to be important for 30 viral propagation. Functions associated with the E4 proteins include DNA replication, late gene

expression, and host cell shutoff. The late gene products include most of the virion capsid proteins, and these are expressed only after most of the processing of a single primary transcript from the major late promoter has occurred. The major late promoter (MLP) exhibits high efficiency during the late phase of the infection (Stratford-Perricaudet and Perricaudet, 1991).

5 As only a small portion of the viral genome appears to be required *in cis* (Tooze, 1981), adenovirus-derived vectors offer excellent potential for the substitution of large DNA fragments when used in connection with cell lines such as 293 cells. Ad5-transformed human embryonic kidney cell lines (Graham, *et al.*, 1977) have been developed to provide the essential viral 10 proteins *in trans*. The characteristics of adenoviruses rendered them good candidates for use in gene transfer both *in vitro* and *in vivo* (Grunhaus & Horwitz, 1992).

15 Particular advantages of an adenovirus system for delivering foreign proteins to a cell include (i) the ability to substitute relatively large pieces of viral DNA by foreign DNA; (ii) the structural stability of recombinant adenoviruses; (iii) the safety of adenoviral administration to humans; and (iv) lack of any known association of adenoviral infection with cancer or malignancies; (v) the ability to obtain high titers of the recombinant virus; and (vi) the high 20 infectivity of adenovirus.

25 Further advantages of adenovirus vectors over retroviruses include the higher levels of gene expression. Additionally, adenovirus replication is independent of host gene replication, unlike retroviral sequences. Because adenovirus transforming genes in the E1 region can be readily deleted and still provide efficient expression vectors, oncogenic risk from adenovirus vectors is thought to be negligible (Grunhaus & Horwitz, 1992).

30 In general, adenovirus gene transfer systems are based upon recombinant, engineered adenovirus which is rendered replication-incompetent by deletion of a portion of its genome, such as E1, and yet still retains its competency for infection. Sequences encoding relatively large foreign proteins can be expressed when additional deletions are made in the adenovirus genome. For example, adenoviruses deleted in both E1 and E3 regions are capable of carrying up to 10 kB

of foreign DNA and can be grown to high titers in 293 cells (Stratford-Perricaudet and Perricaudet, 1991). Persistent expression of transgenes following adenoviral infection has also been reported.

iii. Other Vectors as Expression Constructs

Other viral vectors may be employed as expression constructs in the present invention. Vectors derived from viruses such as vaccinia virus (Ridgeway, 1988; Baichwal and Sugden, 1986; Coupar *et al.*, 1988) adeno-associated virus (AAV) (Ridgeway, 1988; Baichwal and Sugden, 1986; Hermonat and Muzychka, 1984) and herpes viruses may be employed. These viruses offer several attractive features for gene transfer into various mammalian cells (Friedmann, 1989; Ridgeway, 1988; Baichwal and Sugden, 1986; Coupar *et al.*, 1988; Horwitz *et al.*, 1990).

With the recent recognition of defective hepatitis B viruses, new insight was gained into the structure-function relationship of different viral sequences. *in vitro* studies showed that the virus could retain the ability for helper-dependent packaging and reverse transcription despite the deletion of up to 80% of its genome (Horwitz *et al.*, 1990). This suggested that large portions of the genome could be replaced with foreign genetic material. The hepatotropism and persistence (integration) were particularly attractive properties for gene transfer into liver cells. Chang *et al.* recently introduced the chloramphenicol acetyltransferase (CAT) gene into duck hepatitis B virus genome in the place of the polymerase, surface, and pre-surface coding sequences. It was cotransfected with wild-type virus into an avian hepatoma cell line. Culture media containing high titers of the recombinant virus were used to infect primary duckling hepatocytes. Stable CAT gene expression was detected for at least 24 days after transfection (Chang *et al.*, 1991).

iv. Alternative Delivery Systems

In order to effect expression of reporter gene constructs, the expression vector must be delivered into a cell. As described above, one mechanism for delivery is via viral infection where the expression vector is encapsidated in an infectious adenovirus particle.

Several non-viral methods for the transfer of expression vectors into cultured mammalian cells also are contemplated by the present invention. These include calcium phosphate precipitation (Graham and Van Der Eb, 1973; Chen and Okayama, 1987; Rippe *et al.*, 1990) DEAE-dextran (Gopal, 1985), electroporation (Tur-Kaspa *et al.*, 1986; Potter *et al.*, 1984), direct 5 microinjection (Harland and Weintraub, 1985), DNA-loaded liposomes (Nicolau and Sene, 1982; Fraley *et al.*, 1979) and lipofectamine-DNA complexes, cell sonication (Fechheimer *et al.*, 1987), gene bombardment using high velocity microprojectiles (Yang *et al.*, 1990), polycations (Boussif *et al.*, 1995) and receptor-mediated transfection (Wu and Wu, 1987; Wu and Wu, 1988).

10 In one embodiment of the invention, the expression vector may simply consist of naked recombinant vector. Transfer of the construct may be performed by any of the methods mentioned above which physically or chemically permeabilize the cell membrane. For example, Dubensky *et al.* (1984) successfully injected polyomavirus DNA in the form of CaPO₄ precipitates into liver and spleen of adult and newborn mice demonstrating active viral replication and acute infection. Benvenisty and Neshif (1986) also demonstrated that direct 15 intraperitoneal injection of CaPO₄-precipitated plasmids results in expression of the transfected genes. It is envisioned that DNA encoding a construct according to the present invention may also be transferred in a similar manner.

20 Another embodiment of the invention for transferring a naked DNA expression vector into cells may involve particle bombardment. This method depends on the ability to accelerate DNA coated microprojectiles to a high velocity allowing them to pierce cell membranes and enter cells without killing them (Klein *et al.*, 1987). Several devices for accelerating small particles have been developed. One such device relies on a high voltage discharge to generate an 25 electrical current, which in turn provides the motive force (Yang *et al.*, 1990). The microprojectiles used have consisted of biologically inert substances such as tungsten or gold beads.

30 In a further embodiment of the invention, the expression vector may be entrapped in a liposome. Liposomes are vesicular structures characterized by a phospholipid bilayer membrane

and an inner aqueous medium. Multilamellar liposomes have multiple lipid layers separated by aqueous medium. Liposomes form spontaneously when phospholipids are suspended in an excess of aqueous solution. The lipid components undergo self-rearrangement before the formation of closed structures and entrap water and dissolved solutes between the lipid bilayers 5 (Ghosh and Bachhawat, 1991). Also contemplated are lipofectamine-DNA complexes.

Liposome-mediated polynucleotide delivery and expression of foreign DNA *in vitro* has been very successful. Wong *et al.* (1980) demonstrated the feasibility of liposome-mediated delivery and expression of foreign DNA in cultured chick embryo, HeLa and hepatoma cells. 10 Nicolau *et al.* (1987) accomplished successful liposome-mediated gene transfer in rats after intravenous injection.

In certain embodiments of the invention, the liposome may be complexed with a hemagglutinating virus (HVJ). This has been shown to facilitate fusion with the cell membrane and promote cell entry of liposome-encapsulated DNA (Kaneda *et al.*, 1989). In other 15 embodiments, the liposome may be complexed or employed in conjunction with nuclear non-histone chromosomal proteins (HMG-1) (Kato *et al.*, 1991). In yet further embodiments, the liposome may be complexed or employed in conjunction with both HVJ and HMG-1. In that such expression vectors have been successfully employed in transfer and expression of a 20 polynucleotide *in vitro* and *in vivo*, then they are applicable for the present invention. Where a bacteriophage promoter is employed in the DNA construct, it also will be desirable to include within the liposome an appropriate bacteriophage polymerase.

Another mechanism for transferring expression vectors into cells is receptor-mediated 25 delivery. This approach takes advantage of the selective uptake of macromolecules by receptor-mediated endocytosis in almost all eukaryotic cells. Because of the cell type-specific distribution of various receptors, the delivery can be highly specific (Wu and Wu, 1993). Receptor-mediated gene targeting vehicles generally consist of two components: a cell 30 receptor-specific ligand and a DNA-binding agent. Several ligands have been used for receptor-mediated gene transfer. The most extensively characterized ligands are

asialoorosomucoid (ASOR) (Wu and Wu, 1987) and transferrin (Wagner *et al.*, 1993). Recently, a synthetic neoglycoprotein, which recognizes the same receptor as ASOR, has been used as a gene delivery vehicle (Ferkol *et al.*, 1993; Perales *et al.*, 1994) and epidermal growth factor (EGF) has also been used to deliver genes to squamous carcinoma cells (Myers, EPO 0273085).

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In other embodiments, the delivery vehicle may comprise a ligand and a liposome. For example, Nicolau *et al.* (1987) employed lactosyl-ceramide, a galactose-terminal asialganglioside, incorporated into liposomes and observed an increase in the uptake of the insulin gene by hepatocytes. Thus, it is feasible that an adenoviral expression vector also may be 10 specifically delivered into a cell type such as lung, epithelial or tumor cells, by any number of receptor-ligand systems, with or without liposomes. For example, epidermal growth factor (EGF) may be used as the receptor for mediated delivery to cells that exhibit upregulation of EGF receptor, such as tumor cells. Galactose can be used to target the asialoglycoprotein receptor on liver cells. Also, antibodies to CD5 (CLL), CD22 (lymphoma), CD25 (T-cell 15 leukemia) and MAA (melanoma) can similarly be used as targeting moieties.

D. CELL LINES

The cells which can be utilized for transfection in the present invention include and cell that has the transcriptional machinery necessary to direct synthesis from a PTH-rP promoter. Preferred examples include the human cancer cells designated MDA-MB-231, MCF-7, RWGT2 and any other cancer cells which behave as cancer cells *in vivo*, *i.e.*, produce small amounts of PTH-rP unless present in bone. In addition, a number of normal bone cell lines available commercially MC3T3-E1 cells, MG-63 cells, U2OS cells, UMR-106 cells, ROS 17/2.8 cells, SaOS-2 cells, in the catalog provided from the American Type Culture Collection (ATCC), also can be utilized for this assay.

10

E. CANDIDATE COMPOUNDS

Six candidate compounds and their analogs have been identified on the basis of the assay of the present invention. Five of these compounds come from the MicroMolecular Library of MicroSource Discovery Systems, Inc. Gaylordsville, CT - OSW₁ (acriflavinium hydrochloride), OSW₂ (2-(2-carboxypyrid-2-yl)-4-methyl-4-propylimidazolidin-5-one(HCl)), OSW₃ (5-benzylidene hydrazino-1,2,3,4-tetrazole) and OSW₄ (glaucarubolone) and OSW₆ (6-thioguanine). The structures of these compounds are illustrated in FIGs. 6, 2, 1 and 3, and 5, respectively.

20 The sixth compound is dexamethasone (pregna-1,4-diene-3,20-dione,9-fluoro-11,17,21-trihydroxy-16-methyl-(11 β ,16 α). The structure of this compound is illustrated in FIG. 4. Dexamethasone possesses glucocorticoid activity, for which it is used clinically. It is especially useful as an anti-inflammatory and anti-allergic drug. Topically, it is employed in the treatment of glucocorticoid-responsive dermatoses. Systematically, it decreases the incidence and severity 25 of hearing loss subsequent to bacterial meningitis. It also is used as a diagnostic for Cushing's syndrome, lowering plasma cortisol levels in persons that do not have Cushing's. The plasma half-life is 3 to 4 hours, and the biological half-life is 36 to 54 hours.

Dexamethasone is soluble in alcohol (1 g in 42 ml) and practically insoluble in water. Formulation include aerosol (topical), elixir, gel, ophthalmic suspension and tablet. Therapeutic doses range from 500 μ g to 9 mg daily, usually less for maintenance, or 8 mg every other day for one month.

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F. PHARMACEUTICALS AND METHODS OF TREATMENT

In another embodiment of the present invention, there are provided methods for the treatment of cancer. The present invention contemplates the use of compounds having suppressive activity against the PTH-rP promoter and, hence, against PTH-rP in tumor cells.

10 Treatment methods will involve treating an individual with an effective amount of a PTH-rP inhibitory compound. An effective amount is described, generally, as that amount sufficient to detectably and repeatedly reduce the level of PTH-rP in a cell.

15 In an another embodiment of the present invention, there are provided methods for treating bone damage or underdevelopment. The present invention contemplates the use of compounds having stimulatory activity for the PYH-rP promoter and, hence, production of PTH-rP by bone cells. Treatment methods will involve treating an individual with an effective amount of a PTH-rP stimulatory compound. An effective amount is described, generally, as that amount sufficient to detectably and repeatedly increase the level of PTH-rP in a cell.

20 Administration of the compound to patient will follow general protocols for the administration of chemotherapeutics, taking into account the toxicity, if any, of the compound. It is anticipated that the treatment cycles would be repeated as necessary.

25 Where clinical application of a PTH-rP inhibitory compound is contemplated, it will be necessary to prepare the complex as a pharmaceutical composition appropriate for the intended application. Generally this will entail preparing a pharmaceutical composition that is essentially free of pyrogens, as well as any other impurities that could be harmful to humans or animals. One

also will generally desire to employ appropriate salts and buffers to render the complex stable and allow for complex uptake by target cells.

Aqueous compositions of the present invention comprise an effective amount of the compound, dissolved or dispersed in a pharmaceutically acceptable carrier or aqueous medium. Such compositions can also be referred to as inocula. The phrases "pharmaceutically or pharmacologically acceptable" refer to molecular entities and compositions that do not produce an adverse, allergic or other untoward reaction when administered to an animal, or a human, as appropriate. As used herein, "pharmaceutically acceptable carrier" includes any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents and the like. The use of such media and agents for pharmaceutical active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the active ingredient, its use in the therapeutic compositions is contemplated. Supplementary active ingredients also can be incorporated into the compositions.

Solutions of the active compounds as free base or pharmacologically acceptable salts can be prepared in water suitably mixed with a surfactant, such as hydroxypropylcellulose. Dispersions also can be prepared in glycerol, liquid polyethylene glycols, and mixtures thereof and in oils. Under ordinary conditions of storage and use, these preparations contain a preservative to prevent the growth of microorganisms.

The compositions of the present invention may include classic pharmaceutical preparations. Administration of therapeutic compositions according to the present invention will be via any common route so long as the target tissue is available via that route. This includes oral, nasal, buccal, rectal, vaginal or topical. Topical administration would be particularly advantageous for treatment of skin cancers. Alternatively, administration will be by orthotopic, intradermal, subcutaneous, intramuscular, intraperitoneal or intravenous injection. Such compositions would normally be administered as pharmaceutically acceptable compositions that include physiologically acceptable carriers, buffers or other excipients. Intratumoral administration may be preferred for

treatment of discrete tumor masses. Similarly, direct administration into the area of bone damage or underdevelopment may be preferred for treatment of the disorders.

The therapeutic compositions of the present invention are advantageously administered in the form of injectable compositions either as liquid solutions or suspensions; solid forms suitable for solution in, or suspension in, liquid prior to injection may also be prepared. These preparations also may be emulsified. A typical composition for such purpose comprises a pharmaceutically acceptable carrier. For instance, the composition may contain 10 mg, 25 mg, 50 mg or up to about 100 mg of human serum albumin per milliliter of phosphate buffered saline. Other pharmaceutically acceptable carriers include aqueous solutions, non-toxic excipients, including salts, preservatives, buffers and the like. Examples of non-aqueous solvents are propylene glycol, polyethylene glycol, vegetable oil and injectable organic esters such as ethyloleate. Aqueous carriers include water, alcoholic/aqueous solutions, saline solutions, parenteral vehicles such as sodium chloride, Ringer's dextrose, *etc.* Intravenous vehicles include fluid and nutrient replenishers. Preservatives include antimicrobial agents, anti-oxidants, chelating agents and inert gases. The pH and exact concentration of the various components the pharmaceutical composition are adjusted according to well known parameters.

Additional formulations are suitable for oral administration. Oral formulations include such typical excipients as, for example, pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate and the like. The compositions take the form of solutions, suspensions, tablets, pills, capsules, sustained release formulations or powders. When the route is topical, the form may be a cream, ointment, salve or spray.

The term "unit dose" refers to physically discrete units suitable for use in a subject, each unit containing a predetermined-quantity of the therapeutic composition calculated to produce the desired responses, discussed above, in association with its administration, *i.e.*, the appropriate route and treatment regimen. The quantity to be administered, both according to number of treatments and unit dose, depends on the subject to be treated, the state of the subject and the protection

desired. Precise amounts of the therapeutic composition also depend on the judgment of the practitioner and are peculiar to each individual.

Therapies according to the present invention encompass combination therapies that include 5 treatment with anti-PTH-rP compositions as well as standard chemo- and radiotherapies. For example, chemotherapeutics include, but are not limited to, cisplatin (CDDP), carboplatin, procarbazine, mechlorethamine, cyclophosphamide, ifosfamide, melphalan, chlorambucil, bisulfan, nitrosurea, dactinomycin, daunorubicin, doxorubicin, bleomycin, plicomycin, mitomycin, etoposide (VP16), tamoxifen, taxol, transplatin, 5-fluorouracil, vincristin, 10 vinblastin and methotrexate. Also included in combined therapies may be x- and γ -irradiation.

EXAMPLES

The following examples are included to demonstrate preferred embodiments of the 15 invention. It should be appreciated by those of skill in the art that the techniques disclosed in the examples which follow represent techniques discovered by the inventor to function well in the practice of the invention, and thus can be considered to constitute preferred modes for its practice. However, those of skill in the art should, in light of the present disclosure, appreciate that many changes can be made in the specific embodiments which are disclosed and still obtain 20 a like or similar result without departing from the spirit and scope of the invention.

In the examples which follow, the following abbreviations apply: eq (equivalents); M (Molar); mM (millimolar); μ M (micromolar); N (normal); mol (moles); mmol (millimoles); μ mol (micromoles); nmol (nanomoles); kg (kilograms); gm (grams); mg (milligrams); μ g (micrograms); μ l (microliters); vol (volumes); $^{\circ}$ C (degrees Centigrade); FCS (fetal calf serum). 25

Example 1: Transfection of Tumor Cells with PTH-rP promoter linked to a Reporter Gene

To perform studies on the PTH-rP promoter in whole cells, a human breast cancer cell line, MDA-MB-231 (kindly provided by C. Kent Osborne, San Antonio TX) and a human lung cancer cell line, RWGT2, were employed. Various PTH-rP promoters, described in Wysolmerski *et al.*, 1996, were linked with the reporter gene firefly luciferase and stably transfected into cell lines. Plasmid constructs included pGL3B-PTH-rP1.1 and pGL4B-PTHrP 4.0. RWGT2 cells and MDA-MB-231 cells were cultured in DMEM (Life Technologies, Grand Island, NY) with 10% FCS (Hyclone, Logan, UT), 0.1% penicillin/streptomycin and 0.1% nonessential amino acids (Gibco) in 37°C atmosphere of 5% CO₂ /air.

Before stable transfection of MDA-MB-231 cells with pGL3B-PTH-rP1.1, the concentration of geneticin (G418) required to kill MDA 231 cells was determined to be 500 µg/ml. Next, 6-well tissue culture plates were seeded with exponentially growing MDA-MB-231 cells 24 hours prior to transfection. Thirty µg pGL3B-PTH-rP1.1 was diluted with dH₂O to 90 µl and 3 µg pSV2neo added. Ten µl 2.5M CaCl₂ was added followed by 100 µl of 2X BBS. This mixture was next incubated at room temperature for 15 minutes. The mixture was then added to MDA-MB-231 cells dropwise and swirled to distribute evenly, followed by incubation overnight at 37° C in 3% CO₂. The cells were then washed twice with PBS and 3 ml DMEM 10% FCS was added and incubated overnight at 37° C in 5% CO₂. After 3 days, the cells were split 1:10 and incubated for another 3 days at 37° C in 5% CO₂. The cells were washed twice with PBS. DMEM 10% FCS was added containing 500 µg/ml of geneticin. The cells were passaged twice a week for three weeks and incubated at 37° C in 5% CO₂. Subcloning and selection was then begun.

Subcloning consisted of diluting the geneticin resistant MDA-MB-2311 cells containing pGL3B-PTHrP1.1 to a concentration of 0.3 cells/100 µl. Cells were aliquoted in 100 µl increments into ten 96 well tissue culture plates. Colonies were allowed to grow over a period of 3 weeks and then assayed for luciferase activity. The luciferase activity was assessed by

luminometer. Cell lysates were prepared and the luciferase enzyme assay was carried out using assay protocols and the assay kit from Promega (Madison, WI). Routinely, 20 μ l of cell lysate was mixed with 100 μ l of luciferase assay reagent (270 μ M co-enzyme A, 470 μ M luciferin and 530 μ M ATP) and the luciferase activity was measured for 10 seconds in a Turner TD-20e 5 luminometer. The values were normalized with respect to the protein concentration obtained from each experimental sample. The protein concentration was assessed using BioRed reagents. Colonies exhibiting high luciferase activity were further selected by treatment with 2.5 mM dexamethasone. One clone was chosen which exhibited high basal levels of luciferase activity and when dexamethasone was added demonstrated a 30-40% inhibition of luciferase activity. 10 This clone (Clone 30) was used in our screening assay.

Example 2: Analysis of the Ability of Compounds to Inhibit Expression of PTH-rP *In Vitro*

15 To demonstrate that the present invention is useful in evaluating chemical compounds and agents which regulate PTH-rP production, a random array of chemical compounds obtained commercially from a library of 3840 compounds, purchased from MicroSource Inc., (New Haven, CT) was screened. Approximately 342 of 3840 such compounds decreased production of PTH-rP in the present assay system at a concentration of 10 μ M when compared to basal control 20 levels. Such compounds identified from the random library were then tested again at 2 doses (10 μ M, 1 μ M) to again demonstrate that they inhibited PTH-rP promoter-luciferase expression at both doses. These compounds were then tested for toxic effects on MDA-MB-231 cells by the capacity of 90% of the cells or more to exclude trypan blue following 10 minutes incubation with the compounds.

25 The five agents that satisfied these criteria (*i.e.*, effects at both doses, nontoxic to tumor cells), were then tested for their capacity to inhibit PTH-rP production of MDA-MB-231 and RWGT2 cells as measured by RIA. PTH-rP RIA assay was carried out using assay protocols and the PTH-rP assay kit from Nichols Institute Diagnostics (San Juan Capistrano, CA). Routinely, 30 200 μ l of sample was mixed with 100 μ l of the 125 I PTH-RP antibody solution and allowed to

incubate at room temperature for 21 hrs. An avidin coated bead was then added and 90 min later sample/antibody solution mixture was completely aspirated. Beads were washed twice and counted using a Pharmacia Gamma Counter. Two of the 432 compounds were positively identified.

5

Example 3: Analysis of the Ability of Compounds to Inhibit Expression of PTH-rP *In Vivo*. Hypercalcemia Studies

Compounds which are identified as regulating PTH-rP production were initially tested for 10 their capacity to inhibit hypercalcemia *in vivo* by injecting tumor cells into nude mice and treating the mice with the potential inhibitory compounds. Specifically, compounds to be tested were injected into the subcutaneous tissue, intramuscularly, or intravenously into nude mice which were also inoculated with the human cancer cells which produce PTH-rP. The effects on the capacity of the tumor cells to cause hypercalcemia are then assessed over 2-4 weeks.

15 To demonstrate initially that compounds inhibit PTH-rP expression *in vivo*, a human tumor cell which is known to overexpress PTH-rP was used. The tumor, designated RWGT2, is a human squamous cell carcinoma of the lung that was obtained from a hypercalcemic cancer patient (Guise et al, 1993). Metastatic tumor tissue was taken at the time of surgical repair of a pathologic fracture of the femur and implanted into athymic nude mice. The mice developed 20 hypercalcemia, tumors, and increased serum PTH-rP concentrations. Tumor tissue from the mice was used to establish this cell line in culture. The cells produced PTH-rP in culture and reliably produced tumors and hypercalcemia when inoculated intramuscularly into the nude mice.

25 The RWGT2 cells are grown in culture in T75 flasks in α -minimal essential medium (α MEM, Hazelton Biologics Inc., Lenexa, KS) with 10% fetal calf serum and passaged twice a week. The mice are inoculated with RWGT2 cells by subcutaneous inoculation of 1×10^6 cells into the right flank of the upper leg.

30

All animal studies are conducted in accordance with the NIH Guide for the Care and Use of Laboratory Animals. Male BALB/c nude mice (Harlen), 4-6 weeks of age, are housed in a laminar flow isolator hood with a 12-hour-light/12-hour-dark cycle. Water, supplemented with multivitamins (Lymphomed, Melrose Park, IL) and autoclaved mouse chow (Ralston-Purina, St. Louis, MO) were provided *ad libitum*.

Weekly, the mice (each group n=5) are weighed and tumor diameters (long axis and mean axis width) are measured using calipers. Also whole blood samples for ionized calcium (Ca^{2+}) determination is obtained by retro-orbital puncture under anesthesia and measured using a Ciba 10 Corning 634 ISE Ca^{++} /pH analyzer. Finally, blood samples for PTH-rP measurements is obtained by retro-orbital puncture and collected on ice, into pediatric vacutainer tubes containing ethylenediamine tetraacetate (EDTA) (Becton Dickinson, Rutherford, NJ) and aprotinin (Sigma Chemical Company, St. Louis, MO), 400KIU/ml and stored at -70°C. Immunoreactive PTH-rP is measured using a 2 site IRMA kit available from Nichols Institute (San Juan Capistrano, CA).

15 For all experiments compound is administered when hypercalcemia is evident (treatment phase) or from the time of inoculation of RWGT2 cells (prevention phase). OSW3 (FIG. 1) at 50mg/kg/day is able to lower whole blood calcium concentrations (FIG. 7) after hypercalcemia is evident when compared to mice injected with vehicle alone. In addition, OSW3 at 50mg/kg/day is able to lower whole blood calcium concentrations (FIGS. 8A and 8B) and PTH-rP concentrations (FIG. 9) when administered from day 0 when compared to mice injected with vehicle alone.

20 OSW6 (FIG. 5) at 10mg/kg/day is able to lower whole blood calcium concentrations (FIGS. 10A and 10B) after hypercalcemia is evident when compared to mice injected with vehicle alone. The percent change in whole blood calcium levels when compared to mice injected with vehicle alone is significant (FIGS. 10A and 10B). The size of the tumor is also reduced when hypercalcemic mice are treated with OSW6 (FIG. 11). In addition OSW6 at 10mg/kg/day is able to significantly lower whole blood calcium concentrations when 25 administered from day 0 when compared to mice injected with vehicle alone (FIG. 12).

Example 4: Analysis of the Ability of Compounds to Inhibit Expression of PTH-rP *In Vivo*. Bone Metastases Studies.

Compounds which are identified as regulating PTH-rP production were initially tested for 5 their capacity to inhibit bone metastases *in vivo* by injecting tumor cells into nude mice and treating the mice with the potential inhibitory compounds. Specifically, compounds to be tested were injected into the subcutaneous tissue, intramuscularly, or intravenously into nude mice which were also inoculated with the human cancer cells which produce PTH-rP. The effects on the capacity of the tumor cells to cause osteolytic bone lesions are then assessed over 2-4 weeks.

10 To demonstrate that compounds inhibit osteolytic bone metastases *in vivo*, a human tumor cell which is known to express PTH-rP was used. The tumor, named MDA-MB-231, is a human breast cell carcinoma.

15 The MDA-MB-231 cells are grown in culture in T75 flasks in D-minimal essential medium (DMEM, Hazelton Biologics Inc., Lenexa, KS) with 10% fetal calf serum and passaged twice a week. To prepare for mouse inoculation, cells were trypsinized, washed twice with phosphate buffered saline (PBS) and resuspended in PBS to a final concentration of 10^5 cells/100 μ l. The suspended cells are then inoculated into the left ventricle of the heart.

20 25 All animal studies are conducted in accordance with the NIH Guide for the Care and Use of Laboratory Animals. Female BALB/c nude mice (Harlen), 4-6 weeks of age, are housed in a laminar flow isolator hood with a 12-hour-light/12-hour-dark cycle. Water, supplemented with multivitamins (Lymphomed, Melrose Park, IL) and autoclaved mouse chow (Ralston-Purina, St. Louis, MO) were provided *ad libitum*.

30 Weekly, the mice are weighed and whole blood samples for ionized calcium (Ca^{2+}) determination are obtained by retro-orbital puncture under anesthesia and measured using a Ciba Corning 634 ISE Ca^{++} /pH analyzer.

The number of osteolytic bone metastases is determined on radiographs as described by Nakai (1992). Weekly, animals are anesthetized deeply, laid down in a prone position against the film (22x27 cm X-O Mat AR, Kodak, Rochester, NY) and exposed with an x-ray at 35 KVP for 6 seconds using a Faxitron Radiographic Inspection Unit (Model 8050-020, Field Emission Corporation, Inc.). Films are developed using a RP X-O Mat processor (Model M6B, Kodak). The area of osteolytic bone metastases is calculated using a computerized image analysis system in which examination of osteolytic lesions from radiographs is achieved using a fluorescent light box (Kaiser, Germany) and Macro TV Zoom lens 18-108 mm f2,5 (Olympus, Japan) attached to a color video camera (Model DXC-1 51 Sony, Japan). Video images are captured using a frame grabber board (Targa+, Truevision, Inc., USA) with an IBM compatible 486/33Mhz computer system. Quantitation of lesion area is performed using image analysis software (Java, Jandel Video Analysis, Jandel Scientific, CA).

For all experiments compound is administered when osteolytic bone metastases is evident (treatment phase) or from the time of inoculation of MDA-MB-231 cells (prevention phase). OSW3 (FIG. 1) at 50mg/kg/day is able to significantly lower both lesion number and lesion area of osteolytic metastases due to MDA-MB-231 cells when compared with mice injected with vehicle alone (FIGs. 13A and 13B).

Dexamethasone (FIG. 4) at 1 mg/kg/day (low dose) or 2.6 mg/kg/day (high dose) is able to significantly lower both lesion number and lesion area of osteolytic metastases due to MDA-MB-231 cells when compared with mice injected with vehicle alone (FIGs. 14A and 14B).

Example 5: Analysis of the Ability of Compounds to Stimulate Expression of PTH-rP *In Vitro*

Regulation of expression of PTH-rP was assessed by culturing breast cancer cells with and without compounds which might regulate PTH-rP expression. The luciferase activity was assessed by luminometer. Cell lysates were prepared and the luciferase enzyme assay was carried out using assay protocols and the assay kit from Promega (Madison, WI). Routinely, 20

μl of cell lysate was mixed with 100 μl of luciferase assay reagent (270 μM coenzyme A, 470 μM luciferin and 530 μM ATP) and the luciferase activity was measured for 10 seconds in a Turner TD-20e luminometer. The values were normalized with respect to the protein concentration obtained from each experimental sample. The protein concentration was assessed
5 using Bio Rad reagents.

To demonstrate that the present invention is useful in evaluating chemical compounds and agents which regulate PTH-rP production, a random array of chemical compounds obtained commercially from a library of 5000 compounds, purchased from MicroSource Inc., (New
10 Haven, Ct) was screened. Approximately 1 in 2000 such compounds increased production of PYH-rP in the present assay system compared with the positive control. These compounds were shown to be non-toxic for the tumor cells. Such compounds identified from the random library were subjected to detailed dose-response curves to demonstrate that they stimulate PTH-rP messenger RNA expression, and that they increase other effects associated with PTH-rP *in vitro*.
15

The compound designated MS 01500618 was able to stimulate the present assay about 8 fold at a concentration of 3.2 μM (FIGs. 15A and 15B). This compound is acriflavinium hydrochloride, known as OSWs1 (FIG. 6).

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The following references, to the extent that they provide exemplary procedural or other details supplementary to those set forth herein, are specifically incorporated herein by reference.

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SEQUENCE LISTING

(1) GENERAL INFORMATION:

5 (i) APPLICANT: Mundy, Gregory R.
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(ii) TITLE OF INVENTION: SCREENING ASSAY FOR THE IDENTIFICATION OF AGENTS
10 WHICH INHIBIT CANCER METASTASIS TO BONE

10 (iii) NUMBER OF SEQUENCES: 1

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(F) ZIP: 77210

20 (v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
(C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: PatentIn Release #1.0, Version #1.30

25 (vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER: US Unknown
(B) FILING DATE: Concurrently Herewith
(C) CLASSIFICATION: Unknown

30 (viii) ATTORNEY/AGENT INFORMATION:

(A) NAME: Highlander, Steven L.
(B) REGISTRATION NUMBER: 37,642
(C) REFERENCE/DOCKET NUMBER: OSTS:002PZ1

35 (ix) TELECOMMUNICATION INFORMATION:

(A) TELEPHONE: 512/418-3000
(B) TELEFAX: 512/474-7577

40 (2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

45 (A) LENGTH: 4348 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

AGGATCCCAT CAGCTTGATG CATATCTATA CACTCCTCCC TGAGGCAGTT CCTCCAGAGG

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	TTAGCAGCCC	GCCCTGTTCC	TGGAGAAGTC	TTATCCTCAC	CTAACTACAA	AAAGCATTAA	120
	ATAAAGAAAC	ACACCCCTTC	CTATTGTTAC	TTGGGGTTTG	AAGGCATTAC	ATCTTTTTT	180
5	CTTTTTGCT	ACCTTGAGGT	CAGCTGGCAA	CAGCCTCCTC	TCAAGTCTCA	GTCCAGGCGA	240
	GCGATGAGAG	CCACATTCTA	ATGGAATCCA	ATGAGAGCAT	TACTTGAGCT	GATTATGCAA	300
10	CGTCTCTTG	TAAACATTGA	AAAAAGTTG	GAGAAGAGAT	GGGATGAGTT	TAGTTGTTTG	360
	GTTCCAGGGG	ATTTAGAGA	CATACGTTGC	AGCTACAGAT	TGGTAAATGT	GAAATCTGGA	420
	TGCTTATTGG	TAATAAGAGA	ATTCCAAGC	CCAGGTGCAC	TGTTTAAAGT	GCTATAGATT	480
15	CATATTTGGT	TTATAATGTA	TATCTGTTG	CTGTTGGGT	TAAGGAGGAA	AGAAGAAAGA	540
	AGGATGAAGA	GGCTAACGAT	AAATGCTATT	TACTTTTTC	TAAGCTATGA	CAGGAGATAT	600
20	ACATTAAC TG	GGTATTCAAC	TGAATTAAAG	AGTAATGCAT	TTAAAATTT	TTCAACCTCT	660
	ATTAAAATTT	GATATACTGT	AATAAACTGC	CACTGGGGAT	GGGAAGATGG	AAGCCTGGTG	720
	CTCATGGGCT	AGGCATTGT	GTAGGTGTAG	ATCTTATAAT	GCTAAACATG	GAAATACTTC	780
25	AGATTAGAGG	CAGGCCTCCC	ATTTGCTAAG	GTGCATTTAC	ATGACAGCAA	GGCCTAACGCA	840
	AACATTTAGC	TTCTATTGGC	ACTGTTCTA	TTTCTAAACC	TTAGAAAAAA	GGTGTGTGTG	900
30	TGGGCTGGGG	GGACGGGGGG	GGGAGGGAGG	TGGTAGGGGG	TGCTCTGCT	GTGTCTCATT	960
	TGCAGTCATG	CATCCTCTGC	ATTATTATGA	TGGAGATTAC	TCAGTTATGT	TAGGAACTAG	1020
	ATTATGATGT	CAGAAAATAT	CCTTCCAAA	ACAGGCAAAA	AGTCAGGGTC	CTGGGTATAT	1080
35	ATTGAGGAGA	ATTCTACGG	AAAGTAATAA	ACAGGGCAGC	TTGGAAGAGG	TACCTGCTTT	1140
	CTAATAATTG	CCTTAGTGG	GAACAGAAAGT	CTCCTTCAA	GAAGCTTTA	ATTCAATTAA	1200
	AGATTATTT	TTATGTTTC	TGAAGAACAA	CAAAAAATAT	TTCTGGAAAA	GACTGCTGAG	1260
40	AAGATTCCC	TCTTCAGCC	AGAAGAGCAG	AGAGAAGATT	GTAAATCAAG	GAAAAGGTGA	1320
	AGTAATAAAAT	TAGGAGGGAA	CTTGGTATT	CCGAGTATAT	AAAGACTATT	TATTTTCCT	1380
45	GTGTCTATAT	TTTCTCTTTT	TGTGGAGGAG	AGGAAATTCT	AAAAATATT	GATAGATGTT	1440
	TTGCCATTAA	CACCAGAAAA	GTGTGTGGGG	AAAAAGAAAG	GAGGGAAGGA	GTGGGGGGTT	1500
	AATTTGTTT	AATTAGTAGA	AAAAGCAACA	TAAATCAAAG	CAGTCTATTG	ATGCCAGTCC	1560
50	TTAATTTATA	ATGTTCTGAA	AGTAAAGTGA	ATTTATTTAC	AACATAAGTG	ATTTGATAAT	1620
	TTCAATTGAA	TTTTGTTTT	AACCTTCTAT	TGGGAGAAGG	GTTGACTTT	TAAAGCCTGG	1680

	ATAGTTGAA ACTTGGCTAG GTACCTGGA CTTTTTATTG TGGAAGCAAA TATTATCATT	1740
	TCAATGTTAA ACAACTTGCA AGTATTAAAT GGCTCATTG TGATTGACTT TTTTTTTTT	1800
5	TTTTTTTTT TTTTACAGAT TTCCCCCTTC AGATCTAACG ATTACATTAG GGCTCCTGCA	1860
	TCTTTTGGA AGGATTCTTT TTATAAATCA GAAAGTGTTC GAGGTTCAAA GGTTGACATT	1920
10	TCTGAGTGCT GATACTTTGT CCTTTCATAC TATCCAAACA AGTCTAACAT TTAGAAATCC	1980
	TTACACATTC AAGGAAAGTT GTGGAAATTG CCAGAGAGAG AGTGTGTGTG TGTGCGTGTG	2040
	TGTGTTTGG TTTTGTGTTG CTTTTTTCT TTGTTAGTGA GAAGAAGCCG AGTCTTTAA	2100
15	GGTACGGGGT TTACAGTAAT GAACTGAGGA AGGCAGGAGG CTTCTAAGAA AATATGCC	2160
	CCCACCCAAC CTAAGCAGTA AACTTTAATC GCTAGCTAGC TGCAGTGAAC CAGTGGGAGC	2220
20	CCCGATGAGC GAGGGTCTCG GTGACAGCGT GCTATTCTC CCACCCCTGGG TAAAATATGT	2280
	GGAGCATCAC CCGGAAAGTC GGGCTTGATA AAGGCCACAT TCCTTGAATC ATCTCAAGAA	2340
	TCTAAATCAC ACTAGCCTTC TAGAAACTAA TGAACCCCTAC CAGCAGGATT GCCTAGAAGA	2400
25	CAAATATCCC TTGAATGGTT CCCAGTCCAC TCGCGCTCTT TTCAAAAAGT TAGAGGAGCC	2460
	CTGGGGAGGG TATCCACTCC CGCTGCAATC CTTCTCTAGA TGATACTACC CAGTAATTCC	2520
	GAGCAGTCTT TCTTCCCCGC CCATTAGCTT TGGAAAGAAC CTCGGCTTTC CCGTCGCTTC	2580
30	TCCCAGGCAG AGCAGCACAT AACCATAGTT CCACTGCATC TGTCCGCTGG CTGCAGCGAC	2640
	TCGGATACAG TCTTCCAAGA ATCTGTAACC TGGGACTTTT GAGGGGGAGG GGACAAGCAG	2700
35	GTAGGGTATC AGAGAAAGGA TGGGTTAGAC TCCCGACCAC GAGTGAAAAG GGCGTGTGC	2760
	GTGCTCCAGG AGTGTGGTC CCCCTCTGCA ATTCAAAAGG GGGATCTCTC CTGTGCGCGG	2820
	GTTTTTGGG ACCGGCTCCA GATGTCTCCC AGCAGTTCTG AACAGCAAA AAGTGCAATT	2880
40	TAGATATGAA ATCTGGAACG GTTTTGTTTC TTCTAAGCAA AAGATCTCCC TCTCTCTAGC	2940
	CGATGCTCCC CACTCAGTTC ATCCCGGGAA TGGGCCAGGG AGGAAGGTTG TCATGCATCG	3000
45	CCCCGAGCTG CCAGGGGAGC TTCGGGCTCC TTAAATTACAC AGGCCAACAG CCCGCGTCCT	3060
	CTCCGCGCAG GCTCCCGTT GCCCGCGGTC CCCGGCCAG CTCCTGGCC TCCTCCTCGT	3120
	CGGTCCGCCCGG CTGGTGGTCT TGGCGCCCGC TCGTCCAGCT CGGCGCGCCG GGGACCGCCG	3180
50	GCTGCCCGGG GCAGTCCGCA CGCCCTCGGG GATCTCGGCT CCCGGATCCG CCGCGCCGGC	3240
	AGGAGCCGGC CGGGCCTGGA GGGAGCAAGC GGATGCGCCC ACGCCCCCGG CACGGGGATG	3300

	GGCGGACAGG	GCCCCGGGCTC	CGGGGTGGGG	CTCGGCAGAG	CTCCTGACAG	CTCCGGGGCT	3360
	CGGCAGCGCG	GGAGGGGGGA	GCTCCGCCGC	TCGCCGCTCA	TTCCCGGCTC	GGGGCTCCCC	3420
5	TCCACTCGCT	CGGGCGGCCGC	GGGGCCCGTT	CGGGCCGCC	GTCGCCGCC	CCGCCCCCG	3480
	CGCGCCCGCC	CGCCAGCCCG	CCTGCGCCCT	CGCTCGCCCC	GCGCGCGTTC	CTAGGGCGCC	3540
10	ACCTCTTGC	GACTAGCTCA	CTTCTCCGGC	AGGTTTGCCT	CGGAGCGTGT	GAACATTCT	3600
	CCGCTCGGTT	TTCAACTCGC	CTCCAACCTG	CGCCGCCCGG	CCAGCATGTC	TCCCCGCCCG	3660
	TGAAGCGGGC	TGCCGCCTCC	CTGCCGCTCC	GGCTGCCACT	AACGACCCGC	CCTCGCCGCC	3720
15	ACCTGGCCCT	CCTGATCGAC	GACACACGCA	CTTGAAACTT	GTTCTCAGGG	TGTGTGGAAT	3780
	CAACTTTCCG	GAAGCAACCA	GCCCACCAGA	GGAGGTAGAC	AGACAGCTAT	GTATATATAT	3840
20	GTGGGTTTCG	CTACAAGTGG	CTCTGGAACG	AAAGGGCCTG	GTTCGCAAAG	AAGCTGACTT	3900
	CAGAGGGGGGA	AACTTTCTTC	TTTTAGGAGG	CGGTTAGCCC	TGTTCCACGA	ACCCAGGAGA	3960
	ACTGCTGGCC	AGATTAATTA	GACATTGCTA	TGGGAGACGT	GTAAACACAC	TACTTATCAT	4020
25	TGATGCATAT	ATAAAACCAT	TTTATTTTCG	CTATTATTC	AGAGGAAGCG	CCTCTGATT	4080
	GTTCCTTTTT	TCCCTTTTG	CTCTTTCTGG	CTGTGTGGTT	TGGAGAAAGC	ACAGTTGGAG	4140
	TAGCCGGTTG	CTAAATAAGT	AAGTGCTGAG	AGGCTCCAGA	GAAATTTTT	TTCTTTCAA	4200
30	CTTGGGAGAT	GCCCTTGATG	TTGAAGAGGC	TTTTGAGAG	CGGGCTAAA	AGGGGGAGCG	4260
	GAGTAGTGC	GGGAGATGGA	GAGTCCTGAC	TGACACCTCG	GGTCCCATTC	CCTCTGTTG	4320
35	CAGGTCCCCGA	GCGCGAGCGG	AGACGATG				4348